



Research Advisory Panel of California  
Office of the Attorney General  
455 Golden Gate Avenue, Suite 11000  
San Francisco, CA 94102-7004

For RAPC Office Use Only:

Date Received \_\_\_\_\_

PR# \_\_\_\_\_

## Application for Review

### HUMAN RESEARCH SCHEDULE I OR SCHEDULE II CONTROLLED SUBSTANCES

#### Research Advisory Panel of California

All applicable sections of the application must be completed within the form field provided. Please type or print legibly. Note that certain fields require supporting attachments. Incomplete fields or missing attachments will delay the application process.

#### A. TITLE AND DESCRIPTION OF STUDY

☐ Copy of Study Protocol  
Attached (Required)

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#### B. PRINCIPAL INVESTIGATOR

☐ Copies of CV's of Principal  
Investigator and Sub  
Investigators (Required)

Name: \_\_\_\_\_

Institution: \_\_\_\_\_

Address: \_\_\_\_\_

City, State, Zip: \_\_\_\_\_

Direct Contact Phone Number: \_\_\_\_\_

### C. LOCATION WHERE STUDY WILL BE CONDUCTED

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### D. STUDY AND COMPARATOR DRUGS

(List study and comparator drugs and dosages - attach monograph for each.  
Include placebo if applicable)

Study Drug	Dose Range(s)

### E. SOURCE OF STUDY DRUGS

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### F. PLAN FOR STORAGE AND ACCOUNTABILITY OF STUDY DRUGS

If pharmacy based - storage and accountability plan not required - provide name  
and location of pharmacy

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**G. PLANNED NUMBER OF SUBJECTS**

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**H. STUDY DURATION FOR EACH SUBJECT**

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**I. ANTICIPATED STUDY START-UP AND COMPLETION DATES**

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**J. SOURCE OF FUNDING**

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**K. CONSENT**    ☐ Copy Attached of Informed Consent Form to be used  
with Study (Required)

**L. NAME AND ADDRESS OF IRB; IRB REVIEW STATUS**

- ☐ IRB Approval Pending
- ☐ IRB Approval Obtained (Copy Attached)

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**M. *If Applicable* - PROVISIONS FOR MEDICAL EMERGENCIES**

If study drug is being administered at an onsite research lab, office, clinic, or hospital setting, a description of provisions for handling any medical emergencies that might occur is required. Attach description.

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**N. *If Applicable* - PROVISIONS FOR TAKE HOME STUDY MEDS**

If study requires subjects to "take home" single or multiple doses of study meds, a description of provisions for the dispensing and labeling of these medications is required. Attach description. If pharmacy based - dispensing and labeling description not required - provide name and location of pharmacy.

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**O. ACKNOWLEDGMENT & SIGNATURE OF PRINCIPAL INVESTIGATOR**

As a final step in completing this application, Principal Investigator acknowledges that, upon receipt of Panel approval, he/she will comply with all Panel requirements.

\_\_\_\_\_  
Signature

\_\_\_\_\_  
Date